Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID: SSPTAJRK1626

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

```
Web Page URLs for STN Seminar Schedule - N. America
NEWS
     1
                 "Ask CAS" for self-help around the clock
NEWS
     2
NEWS
     3
        DEC 05
                CASREACT(R) - Over 10 million reactions available
                 2006 MeSH terms loaded in MEDLINE/LMEDLINE
NEWS
        DEC 14
                2006 MeSH terms loaded for MEDLINE file segment of TOXCENTER
NEWS
     5
        DEC 14
NEWS
     6
        DEC 14
                CA/CAplus to be enhanced with updated IPC codes
NEWS
        DEC 21
                IPC search and display fields enhanced in CA/CAplus with the
                 IPC reform
NEWS 8
        DEC 23
                New IPC8 SEARCH, DISPLAY, and SELECT fields in USPATFULL/
                 USPAT2
                 IPC 8 searching in IFIPAT, IFIUDB, and IFICDB
NEWS 9
        JAN 13
                New IPC 8 SEARCH, DISPLAY, and SELECT enhancements added to
NEWS 10
        JAN 13
                 INPADOC
NEWS 11
        JAN 17
                 Pre-1988 INPI data added to MARPAT
        JAN 17 IPC 8 in the WPI family of databases including WPIFV
NEWS 12
```

NEWS EXPRESS JANUARY 03 CURRENT VERSION FOR WINDOWS IS V8.01,
CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
AND CURRENT DISCOVER FILE IS DATED 19 DECEMBER 2005.
V8.0 USERS CAN OBTAIN THE UPGRADE TO V8.01 AT
http://download.cas.org/express/v8.0-Discover/

NEWS HOURS STN Operating Hours Plus Help Desk Availability
NEWS INTER General Internet Information
NEWS LOGIN Welcome Banner and News Items
NEWS PHONE Direct Dial and Telecommunication Network Access to STN
NEWS WWW CAS World Wide Web Site (general information)

Enter NEWS followed by the item number or name to see news on that specific topic.

All use of STN is subject to the provisions of the STN Customer agreement. Please note that this agreement limits use to scientific research. Use for software development or design or implementation of commercial gateways or other similar uses is prohibited and may result in loss of user privileges and other penalties.

FILE 'HOME' ENTERED AT 14:13:18 ON 20 JAN 2006

=> file reg
COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION

FULL ESTIMATED COST

0.21 0.21

FILE 'REGISTRY' ENTERED AT 14:13:25 ON 20 JAN 2006 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2006 American Chemical Society (ACS)

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 18 JAN 2006 HIGHEST RN 872163-75-2 DICTIONARY FILE UPDATES: 18 JAN 2006 HIGHEST RN 872163-75-2

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH JULY 14, 2005

Please note that search-term pricing does apply when conducting SmartSELECT searches.

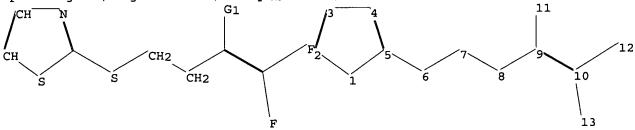
************ The CA roles and document type information have been removed from * the IDE default display format and the ED field has been added, effective March 20, 2005. A new display format, IDERL, is now available and contains the CA role and document type information.

Structure search iteration limits have been increased. See HELP SLIMITS for details.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/ONLINE/UG/regprops.html

=> Uploading C:\Program Files\Stnexp\Queries\10501115\Struc 2.str



chain nodes : 6 7 8 9 10 11 12 13 ring nodes : 1 2 3 4 5 chain bonds : 5-6 6-7 7-8

8-9 9-10 9-11 10-12 10-13

ring bonds :

1-2 1-5 2-3 3-4 4 - 5

exact/norm bonds :

1-2 1-5 2-3 3-4 4-5 5-6 9-11

exact bonds :

6-7 7-8 8-9 9-10 10-12 10-13

G1:H,F

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:CLASS 7:CLASS 8:CLASS 9:CLASS

10:CLASS 11:CLASS 12:CLASS 13:CLASS

L1 STRUCTURE UPLOADED

=> d

L1 HAS NO ANSWERS

L1 STR

G1 H,F

Structure attributes must be viewed using STN Express query preparation.

=> 11

SAMPLE SEARCH INITIATED 14:13:44 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 28 TO ITERATE

100.0% PROCESSED 28 ITERATIONS 0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 243 TO 877

PROJECTED ANSWERS: 0 TO

L2 0 SEA SSS SAM L1

=> 11 full

FULL SEARCH INITIATED 14:13:48 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 598 TO ITERATE

100.0% PROCESSED 598 ITERATIONS 4 ANSWERS

SEARCH TIME: 00.00.01

L3 4 SEA SSS FUL L1

=> d scan

4 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN L3

Thiazole, 2-[(4,4-difluoro-3-butenyl)sulfonyl]- (9CI) IN

MF C7 H7 F2 N O2 S2

$$\begin{array}{c|c}
 & O \\
 & \parallel \\
 & S - CH_2 - CH_2 - CH = CF_2
\end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):3

4 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN L3

IN Thiazole, 2-[(3,4,4-trifluoro-3-butenyl)thio]- (9CI)

C7 H6 F3 N S2

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

4 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN L3

Thiazole, 2-[(4,4-difluoro-3-butenyl)thio]- (9CI) IN Thiazole, 2-[
MF C7 H7 F2 N S2

$$S-CH_2-CH_2-CH=CF_2$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

4 ANSWERS REGISTRY COPYRIGHT 2006 ACS on STN L3

Thiazole, 2-[(4,4-difluoro-3-butenyl)sulfinyl]- (9CI) IN

C7 H7 F2 N O S2 MF

$$\begin{array}{c|c}
 & \circ \\
 & \circ \\$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

ALL ANSWERS HAVE BEEN SCANNED

=> d rn 1-4

L3 ANSWER 1 OF 4 REGISTRY COPYRIGHT 2006 ACS on STN

RN 172933-36-7 REGISTRY

L3 ANSWER 2 OF 4 REGISTRY COPYRIGHT 2006 ACS on STN

RN 172933-35-6 REGISTRY

L3 ANSWER 3 OF 4 REGISTRY COPYRIGHT 2006 ACS on STN

RN 172933-05-0 REGISTRY

L3 ANSWER 4 OF 4 REGISTRY COPYRIGHT 2006 ACS on STN

RN 109993-23-9 REGISTRY

=> d hitstr 1-4
'HITSTR' IS NOT A VALID FORMAT FOR FILE 'REGISTRY'

The following are valid formats:

Substance information can be displayed by requesting individual fields or predefined formats. The predefined substance formats are: (RN = CAS Registry Number)

REG - RN

SAM - Index Name, MF, and structure - no RN

FIDE - All substance data, except sequence data

IDE - FIDE, but only 50 names
SQIDE - IDE, plus sequence data

SQIDE3 - Same as SQIDE, but 3-letter amino acid codes are used

SQD - Protein sequence data, includes RN

SQD3 - Same as SQD, but 3-letter amino acid codes are used

SQN - Protein sequence name information, includes RN

CALC - Table of calculated properties EPROP - Table of experimental properties

PROP - EPROP and CALC

Any CA File format may be combined with any substance format to obtain CA references citing the substance. The substance formats must be cited first. The CA File predefined formats are:

ABS -- Abstract

APPS -- Application and Priority Information

BIB -- CA Accession Number, plus Bibliographic Data

CAN -- CA Accession Number

CBIB -- CA Accession Number, plus Bibliographic Data (compressed)

IND -- Index Data

IPC -- International Patent Classification

PATS -- PI, SO

STD -- BIB, IPC, and NCL

IABS -- ABS, indented, with text labels IBIB -- BIB, indented, with text labels

ISTD -- STD format, indented

OBIB ----- AN, plus Bibliographic Data (original)

OIBIB ----- OBIB, indented with text labels

SBIB ----- BIB, no citations SIBIB ----- IBIB, no citations

The ALL format gives FIDE BIB ABS IND RE, plus sequence data when it is available.

The MAX format is the same as ALL.

The IALL format is the same as ALL with BIB ABS and IND indented, with text labels.

For additional information, please consult the following help messages:

HELP DFIELDS -- To see a complete list of individual display fields. HELP FORMATS -- To see detailed descriptions of the predefined formats. ENTER DISPLAY FORMAT (IDE):sam

L3 ANSWER 1 OF 4 REGISTRY COPYRIGHT 2006 ACS on STN

IN Thiazole, 2-[(4,4-difluoro-3-butenyl)sulfonyl]- (9CI)

MF C7 H7 F2 N O2 S2

$$\begin{array}{c|c}
 & O \\
 & \parallel \\
 & S - CH_2 - CH_2 - CH = CF_2
\end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L3 ANSWER 2 OF 4 REGISTRY COPYRIGHT 2006 ACS on STN

IN Thiazole, 2-[(4,4-difluoro-3-butenyl)sulfinyl]- (9CI)

MF C7 H7 F2 N O S2

$$\begin{array}{c|c}
\circ \\
\parallel \\
s-cH_2-CH_2-CH_2-CH_2
\end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L3 ANSWER 3 OF 4 REGISTRY COPYRIGHT 2006 ACS on STN IN Thiazole, 2-[(4,4-difluoro-3-butenyl)thio]- (9CI) MF C7 H7 F2 N S2

$$S - CH_2 - CH_2 - CH = CF_2$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L3 ANSWER 4 OF 4 REGISTRY COPYRIGHT 2006 ACS on STN IN Thiazole, 2-[(3,4,4-trifluoro-3-butenyl)thio]- (9CI) MF C7 H6 F3 N S2

$$S = CH_2 - CH_2 - C = F$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

=> file caplus
COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 173.30 173.51

FULL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 14:16:22 ON 20 JAN 2006 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS)

Copyright of the articles to which records in this database refer is held by the publishers listed in the PUBLISHER (PB) field (available for records published or updated in Chemical Abstracts after December 26, 1996), unless otherwise indicated in the original publications. The CA Lexicon is the copyrighted intellectual property of the American Chemical Society and is provided to assist you in searching databases on STN. Any dissemination, distribution, copying, or storing of this information, without the prior written consent of CAS, is strictly prohibited.

FILE COVERS 1907 - 20 Jan 2006 VOL 144 ISS 5 FILE LAST UPDATED: 19 Jan 2006 (20060119/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

http://www.cas.org/infopolicy.html

=> 172933-05-0

REG1stRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress... Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L5 1 L4

=> 15

L6 1 L4

=> d ibib abs hitstr

L6 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

1995:994834 CAPLUS

DOCUMENT NUMBER:

124:117350

TITLE:

Preparation of (4,4-difluorobut-3-enylthio)-

substituted heterocyclic or carbocyclic ring compounds

having pesticidal activity

INVENTOR(S):

Turnbull, Michael Drysdale; Bansal, Harjinder Singh; Smith, Alison Mary; Salmon, Roger; Fitzjohn, Steven; Godrey, Christopher Richard Ayles; Hotson, Matthew Brian; Sillars, Nan Catherine; Dowling, Alan John

PATENT ASSIGNEE(S):

SOURCE:

Zeneca Ltd., UK PCT Int. Appl., 194 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

LANGUAGE:

Patent English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PAT	CENT	NO.			KIN	D :	DATE			APPL	ICAT	ION :	NO.		D	ATE		
WO 9524403				A1		1995	0914	,	WO 1	 995-	GB40	0		1:	19950227			
	W:	AM,	AU,	BB,	BG,	BR,	BY,	CA,	CN,	CZ,	EE,	FI,	GE,	HU,	JP,	KE,	KG,	
		KP,	KR,	ΚZ,	LK,	LR,	LT,	LV,	MD,	MG,	MN,	MW,	MX,	NO,	ΝŻ,	PL,	RO,	
		RU,	SD,	SG,	SI,	SK,	ТJ,	TT,	UA,	US,	UΖ,	VN						
	RW:	KE,	MW,	SD,	SZ,	UG,	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙE,	IT,	
		LU,	MC,	NL,	PT,	SE,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	ML,	MR,	ΝE,	
		SN,	TD,	TG														
CA	2182	520			AA		1995	0914	1	CA 1	995-	2182	520		1:	9950	227	
AU	9518	164			A1		1995	0925		AU 1	995-	1816	4		1:	9950	227	
ΑU	6852	42			B2		1998	0115										
ΕP	7494	33			A1		1996:	1227		EP 1	995-	9098	54		1:	9950	227	
\mathbf{EP}	7494	33			B1		2003	0507										
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙE,	IT,	LI,	LU,	MC,	ΝL,	PT,	SE
CN	1143				Α		1997	0226		CN 1	995-	1920	29		1:	9950	227	
	7490				A2		1997	0228		HU 1	996-	2417			1	9950	227	
HU	2152	11			В		1998:	1028										
BR	9507	042			Α		1997	0909				7042				9950		
JP	0951	0197			T2		1997	1014				5232				9950		
CZ	2856				В6		1999	0915				2632				9950		
RU	2151	147			Cl		2000	0620		RU 1	996-	1201	48		1	9950	227	

RO 116399	B1	20010130	RO	1996-1788		19950227
SK 281491	В6	20010409	SK	1996-1148		19950227
AT 239714	É	20030515	AT	1995-909854		19950227
PT 749433	T	20030829	PT	1995-909854		19950227
ES 2199240	Т3	20040216	ES	1995-909854		19950227
US 5705516	Α	19980106	US	1995-400912		19950308
US 5912243	Α	19990615	US	1996-702623		19960828
FI 9603539	Α	19960909	FI	1996-3539		19960909
NO 9603776	Α	19961107	NO	1996-3776		19960909
LV 11686	В	19970620	LV	1996-363		19960910
US 5952359	A	19990914	US	1997-887858		19970703
PRIORITY APPLN. INFO.:			GB	1994-4716	Α	19940310
			GB	1994-4717	Α	19940310
			GB	1994-4718	Α	19940310
			GB	1994-4719	Α	19940310
			GB	1994-4720	Α	19940310
			GB	1994-4721	Α	19940310
			GB	1995-521	Α	19950111
			WO	1995-GB400	W	19950227
			US	1995-400912	A3	19950308
OTHER COIDCE(C).	ת אם מוא או	124.117250				

OTHER SOURCE(S): MARPAT 124:117350

GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

The title compds. represented by the general formula RS(O)nCH2CH2CH:CF2 [n = 0,1,2; R is a group of formulas Q - Q13 (X = O, S), etc., wherein the S(O)mCH2CH2CH:CF2 group is at least one of R1 (when attached to a carbon atom), R2, R3, R4, R5 or R6; e.g. when R1 is attached to a carbon atom, R2, R3, R4, R5 and R6 are each independently H, optionally substituted alkyl, optionally substituted alkenyl, alkynyl, cycloalkyl, alkylcycloalkyl, alkoxy, alkenyloxy, alkynyloxy, hydroxyalkyl, alkoxyalkyl, optionally substituted aryl, optionally substituted arylalkyl, optionally substituted heteroaryl, optionally substituted heteroarylalkyl, optionally substituted aryloxy, optionally substituted arylalkoxy, optionally substituted aryloxyalkyl, optionally substituted heteroaryloxy, optionally substituted heteroarylalkoxy, optionally substituted heteroaryloxyalkyl, haloalkyl, haloalkenyl, haloalkynyl, haloalkoxy, haloalkenyloxy, haloalkynyloxy, halo, HO, cyano, NO2, NR7R8, NR7COR8, NR7CSR8, NR7SO2R8, N(SO2R7)(SO2R8), COR7, CONR7R8, alkyl-CONR7R8, CR7NR8, CO2R7, O2CR7, SR7, SOR7, SO2R7, alkyl-SR7, alkyl-SOR7, alkyl-SO2R7, OSO2R7, SO2NR7R8, CSNR7R9, SiR7R8R9, OCH2CO2R7, OCH2CH2CO2R7, CONR7SO2R8, alkyl-CONR7SO2R8, NHCONR7R8, NHCSNR7R8, or an adjacent pair of R1 - R6 when taken together form a fused 5- or 6-membered carbocyclic or heterocyclic ring] are prepared Thus, a solution of 4,4-difluorobut-3-enyl thioacetate in 50% aqueous NaOH was stirred vigorously for 30 min, followed by successively adding Et 5-chloro-4-methylisoxazole in CH2Cl2 and Bu4NBr, and the reaction mixture was stirred at the ambient temperature for 3 h to give Et

5-(4,4-difluorobut-3-enylthio)-3-methylisoxazole-4-carboxylate. The latter compound was saponified with a mixture of 2 M NaOH and isopropanol and acidified with 2 M HCl to give the acid 5-(4,4-difluorobut-3-enylthio)-3-methylisoxazole-4-carboxylic acid, which was treated with Et chloroformate and Et3N in CH2Cl2 at 0° and then with NH3(g) to give the amide 5-(4,4-difluorobut-3-enylthio)-3-methylisoxazole-4-carboxamide (I). I controlled 100% Tetranychus urticae (spider mite) and Myzus persicae (green peach aphid) upon contract at 100 ppm and 100% Meloidogyne

incognita (root knot nematode) at 2 ppm as a drench solution to 2 wk old cucumber plants.

ΙT 172933-05-0P

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of (difluorobutenylthio)-substituted heterocyclic or carbocyclic ring compds. as pesticides)

RN 172933-05-0 CAPLUS

Thiazole, 2-[(4,4-difluoro-3-butenyl)thio]- (9CI) (CA INDEX NAME) CN

$$S-CH_2-CH_2-CH=CF_2$$

=> 109993-23-9

REG1stRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress... Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

L8 4 L7

=> d ibib abs hitstr

ANSWER 1 OF 4 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

2003:570971 CAPLUS

DOCUMENT NUMBER:

139:133556

TITLE:

Method for producing halogenated 2-(3-butenylthio)-1,3-

thiazoles

INVENTOR(S):

Straub, Alexander

PATENT ASSIGNEE(S):

Bayer CropScience AG, Germany

SOURCE:

PCT Int. Appl., 31 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent German

LANGUAGE:

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.						KIN	ו ס	DATE		1	APPL	ICAT:	ION I	NO.		D	ATE	030103 CH, CN, GE, GH, LK, LR, OM, PH, TT, TZ,		
	WO	2003	0598	96		A1	:	2003	0724	1	WO 2	003-1	EP30	0028		2	0030	103		
		W:	ΑE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,		
			CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,		
			GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	ΚP,	KR,	ΚZ,	LC,	LK,	LR,		
			LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	ΜZ,	NO,	NZ,	OM,	PH,		
			PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	TJ,	TM,	TN,	TR,	TT,	TZ,		
			UA,	ŪĠ,	US,	UΖ,	VC,	VN,	YU,	ZA,	ZM,	ZW								
		RW:	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	ŪĠ,	ZM,	ZW,	AM,	ΑZ,	BY,		
			KG,	ΚZ,	MD,	RU,	ТJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,		
			FI.	FR,	GB,	GR,	HU,	IE,	IT,	LU,	MC,	NL,	PT,	SE,	SI,	SK,	TR,	BF,		

BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG DE 10201238 A1 20030724 DE 2002-10201238 20020115 EP 1467980 A1 20041020 EP 2003-708046 20030103 EP 1467980 **B1** 20050727 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK 20050609 US 2003-501115 20030103 US 2005124816 A1 JP 2005519896 T2 20050707 JP 2003-560000 20030103 AT 300527 E 20050815 AT 2003-708046 20030103 PRIORITY APPLN. INFO.: DE 2002-10201238 20020115 WO 2003-EP28 W 20030103 OTHER SOURCE(S): CASREACT 139:133556; MARPAT 139:133556

GI

Title compds. (I; R = H, F), were prepared in following steps (1) preparing AB F2C:CRCH2CH2SCN (II; R as above) by reacting F2C:CRCH2CH2X (R as above, X = Br, Cl, mesylate, tosylate) with M+SCN-(M+=H, NH+,tetraalkylammonium, alkaline (earth) ion) in the presence of a reaction aid and a solvent, (2) treatment of II with H2S or salts thereof in the presence of a reaction aid and a solvent to give F2C:CRCH2CH2S(:NH)SH (III; R as above), and (3) reacting III with MeCHO, ClCH2CHO, or chloroacetaldehyde dialkylacetal in a solvent to give I. Thus, NH4NCS in EtOH was stirred with 4-bromo-1,1,2-trifluoro-1-butene for 2 h at room temperature to give 93.3% 3,4,4-trifluoro-3-butenylthiocyanate. The latter and Et3N in t-BuOMe were treated with H2S followed by stirring over night at room temperature to give 88.5% 3,4,4-trifluoro-3-butenyldithiocarbamate which was treated with concentrated HCl and 45% ClCH2CHO in dioxane followed by boiling for 4 h whereby ClCH2CHO was again added after 2 h to give 94.4% 2-[(3,4,4-trifluoro-3-butenyl)thio]-1,3-thiazole. I are important intermediates for producing pesticides.

IT 109993-23-9P

> RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)

(method for producing halogenated (butenylthio)thiazoles)

109993-23-9 CAPLUS RN

Thiazole, 2-[(3,4,4-trifluoro-3-butenyl)thio]- (9CI) (CA INDEX NAME) CN

$$S - CH_2 - CH_2 - C - F$$

REFERENCE COUNT: THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS 6 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d ibib abs hitstr 2-4

L8 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2006 ACS on STN

GI

ACCESSION NUMBER: 2003:282551 CAPLUS

DOCUMENT NUMBER: 138:304270

TITLE: Preparation of nematocidal trifluorobutenylthio(or

sulfinyl/sulfonyl) thiazoles

INVENTOR(S): Watanabe, Yukiyoshi; Ishikawa, Koichi; Otsu, Yuich;

Shibuya, Katsuhiko

PATENT ASSIGNEE(S): Bayer CropScience AG, Germany

Ι

SOURCE: PCT Int. Appl., 42 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT	PATENT NO.							7	APPL	ICAT:	ION 1	. O <i>l</i>		DATE					
					-														
WO 2003	WO 2003029231					2003	0410	1	WO 2	002-1	EP10	351		20	20020916				
₩:	AE,	AG,	AL,	AM,	AT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,			
	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,			
	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KΕ,	KG,	ΚP,	KR,	ΚZ,	LC,	LK,	LR,			
	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	ΜZ,	NO,	ΝZ,	OM,	PH,			
	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ΤJ,	TM,	TN,	TR,	TT,	TZ,			
	UA,	UG,	US,	UZ,	VC,	VN,	ΥU,	ZA,	ZM,	zw									
RW:	GH,	GM,	KE,	LS,	MW,	ΜZ,	SD,	SL,	SZ,	TZ,	ŪĠ,	ZM,	ZW,	AM,	ΑZ,	BY,			
	KG,	KZ,	MD,	RU,	TJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,			
	FI,	FR,	GB,	GR,	ΙE,	IT,	LU,	MC,	NL,	PT,	SE,	SK,	TR,	BF,	ВJ,	CF,			
	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG						
JP 2003	A2		2003	0418	,	JP 2	001-	3013	16		20	0010	928						
PRIORITY APP					,	JP 2	001-	3013	16	7	A 20	0010	928						
OTHER SOURCE	MAR	TAS	138:	3042	70														

$$\begin{array}{c|c}
R^1 & F \\
\hline
 & S \\
\hline
 & S \\
\hline
 & S \\
\hline
 & S \\
\hline
 & F
\end{array}$$

AB The title compds. [I; R1 = H, halo, alkyl, haloalkyl, cycloalkyl, alkoxycarbonylmethyl; R2 = H, halo, alkyl, alkoxyalkyl, alkylthioalkyl, carboxy, alkylaminocarbonyl, cycloalkylaminocarbonyl, dialkylaminocarbonyl, alkoxycarbonyl; n = 0-2; with the proviso that R1 and R2 do not represent hydrogen at the same time, and in case R1 represents hydrogen, then R2 does not represent halogen], useful as nematocides, were prepared Thus, reacting 5-ethoxycarbonyl-2-mercapto-4-methylthiazole with 4-bromo-1,1,2-trifluoro-1-butene in the presence of K2CO3 in MeCN afforded 65% I [R1 = Me; R2 = CO2Et; n = 0]. Seven of the prepared compds. I showed more than 90% controlling effect at 10 ppm in test for Meloidogyne spp. (soil pot test).

IT 109993-23-9

RL: RCT (Reactant); RACT (Reactant or reagent)
 (preparation of nematicidal trifluorobutenylthio(or sulfinyl/sulfonyl)
 thiazoles)

RN 109993-23-9 CAPLUS

CN Thiazole, 2-[(3,4,4-trifluoro-3-butenyl)thio]- (9CI) (CA INDEX NAME)

$$S - CH_2 - CH_2 - C - F$$

REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2001:31482 CAPLUS

DOCUMENT NUMBER: 134:100860

TITLE: Nematocidal trifluorobutenes

INVENTOR(S): Watanabe, Yukiyoshi; Ishikawa, Koichi; Otsu, Yuichi;

Shibuya, Katsuhiko; Abe, Takahisa

PATENT ASSIGNEE(S): Nihon Bayer Agrochem K.K., Japan SOURCE: PCT Int. Appl., 27 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.										APPLICATION NO.							DATE			
	WO 2001002378																			
		W:	ΑE,	AG,	AL,	AM,	AT,	AU,	ΑZ,	BA,	BE	, B	3, :	BR,	BY,	ΒZ,	CA,	CH,	CN,	
								DM,												
								JP,												
								MK,												
			SD,	SE,	SG,	SI,	SK,	SL,	TJ,	TM,	TR	, T	Γ,	TZ,	UA,	UG,	US,	UΖ,	VN,	
								BY,												
		RW:						MZ,								AT,	BE,	CH,	CY,	
			DE,	DK,	ES,	FI,	FR,	GB,	GR,	ΙE,	II	', Li	IJ, :	MC,	NL,	PT,	SE,	BF,	ВJ,	
			CF.	CG.	CI.	CM.	GA.	GN.	GW,	ML,	MR	2, NI	E,	SN,	TD,	TG				
	JΡ	2001	0196	85 [.]	•	A2	·	2001	0123		JΡ	1999	9-1	916	38		1	9990	706	
	CA	2378	148			AA		2001	0111		CA	200	0-2	378	148		2	0000	628	
	BR	2000	0122	43		Α		2002	0326		BR	2000	0-1	224	3		2	0000	628	
		1200				A1		2002	0502		ΕP	200	0-9	371	36		2	0000	628	
	ΕP	1200	418																	
		R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GF	, I	Γ,	LI,	LU,	NL,	SE,	MC,	PT,	
								RO,												
	TR	2002	0006	8		T2		2002	0521		TR	200	2-2	002	00068	3	2	0000	628	
	JP	2003	5034	85		T2		2003	0128		JP	200	1-5	078	16		2	0000	628	
	ΑT	2631	57			E		2004	0415		ΑT	200	0-9	371	36		2	0000	628	
	ES	2215	671			Т3		2004	1016		ES	200	0-9	371	36		2	0000		
	ZA	2001	0099	95		A		2002	0827		ZA	200	1-9	995			2	0011	205	
	US	6734	198			В1		2004	0511		US	200	2 - 3	036	1		2	0020	305	
	нк	1046	403			Al		2005	0422		HK	200	2-1	076	54		2	0021	022	
PRIOR	IT	APP	LN.								JP	199	9-1	916	38		A 1	9990	706	
															8			0000		
ОТНЕВ	90	TIRCE	(8) .			MAR	РАТ	134 .	1008	60										

OTHER SOURCE(S): MARPAT 134:100860

GI

$$X \xrightarrow{N} S(0)_{nCH_2CH_2CF=CF_2}$$

AB Title compds. I (n = 0, 1, 2; X = halo) were prepared Thus, 4.8 g N-chlorosuccinimide was added to a solution of 6.75 g 2-[(3,4,4-trifluoro-3-butenyl)thio]thiazole in 60 mL CCl4, and the mixture was refluxed for 18 h to give I (n = 0, X = Cl). Oxidation of this product with m-chloroperoxybenzoic acid and with 31% H2O2 gave I (n = 1, X = Cl) and I (n = 2, X = Cl), resp. I (n = 0, 1, 2; X = Cl) showed 100-71% controlling effect against Meloidogyne incognita on tomatoes.

IT 109993-23-9P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)

(preparation and nematicidal activity of)

RN 109993-23-9 CAPLUS

CN Thiazole, 2-[(3,4,4-trifluoro-3-butenyl)thio]- (9CI) (CA INDEX NAME)

$$\begin{array}{c} CF_2 \\ \parallel \\ S - CH_2 - CH_2 - C - F \end{array}$$

REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1987:496721 CAPLUS

DOCUMENT NUMBER: 107:96721

TITLE: Pesticidal (thiadiazolylthio)trifluorobutene analogs

INVENTOR(S): Cullen, Thomas Gerard; Martinez, Anthony Joseph

PATENT ASSIGNEE(S): FMC Corp., USA

SOURCE: PCT Int. Appl., 102 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND DATE	APPLICATION NO.	DATE
WO 8607590	A1 19861231	WO 1986-US1284	19860612
W: AU, BR, DK,	HU, JP, KR		
RW: CF, CG, CH,	CM, DE, FR, GA,	GB, IT, ML, MR, NL, SN,	TD, TG
AU 8661229	A1 19870113	AU 1986-61229	19860612
AU 601656	B2 19900913		
EP 228447	A1 19870715	EP 1986-904515	19860612
R: CH, DE, FR,	GB, IT, LI, NL		
HU 42424	A2 19870728	HU 1986-3254	19860612
HU 204022	B 19911128		
BR 8606746	A 19871013	BR 1986-6746	19860612
JP 63500037	T2 19880107	JP 1986-503571	19860612
CA 1277668	Al 19901211	CA 1986-511879	19860618

CN 86104207	Α	19870401	CN	1986-104207		19860619
ZA 8604637	Α	19880224	ZA	1986-4637		19860620
DK 8700843	Α	19870219	DK	1987-843		19870219
US 4952580	Α	19900828	US	1988-270903		19881109
PRIORITY APPLN. INFO.:			US	1985-746911	A	19850620
			US	1985-747142	A	19850620
			US	1986-870055	B1	19860603
			WO	1986-US1284	Α	19860612
			US	1988-161575	B2	19880229

GI

F2C:CF(CH2)nZR [n = 1-4; Z = S, O, N, CH2; when Z = S, R = thiazolyl, F2C:CFCH2CH2CCCH2, or (un)substituted thienyl, thianaphthyl, thiazolinyl, thiadiazolyl, and oxadiazolyl; when Z = O, R = COR1 where R1 = perfluoroalkyl, dihydrothiazolylthiomethyl, or (un)substituted Ph, thienyl, furanyl, pyrrolyl; when Z = N, ZR = isothiocyanato, succinimido, or saccharin group; when Z = CH2, R = OH], useful as pesticides, were prepared Refluxing a mixture of 0.08 mol NCN:C(S-K+)2 and 0.08 mol S in MeOH gave 18.1 g thiadiazole derivative I (R2 = R3 = K), which was alkylated by BrCH2CH2CF:CF2 in MeCOEt to give I (R2 = R3 = CH2CH2CF:CF2), which at 5 ppm completely controlled the root-knot nematode.

IT 109993-23-9P

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of, as pesticide)

RN 109993-23-9 CAPLUS

CN Thiazole, 2-[(3,4,4-trifluoro-3-butenyl)thio]- (9CI) (CA INDEX NAME)

$$\begin{array}{c} \begin{array}{c} CF_2 \\ \parallel \\ S \end{array} \\ S - CH_2 - CH_2 - C - F \end{array}$$

=> file caplus scisearch medline COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION 25.15 208.44 FULL ESTIMATED COST DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL SESSION ENTRY -3.75 CA SUBSCRIBER PRICE -3.00

FILE 'CAPLUS' ENTERED AT 14:22:13 ON 20 JAN 2006 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'SCISEARCH' ENTERED AT 14:22:13 ON 20 JAN 2006 Copyright (c) 2006 The Thomson Corporation

FILE 'MEDLINE' ENTERED AT 14:22:13 ON 20 JAN 2006

=> straub

L9 1096 STRAUB

=> 19 and alexander

L10 0 L9 AND ALEXANDER

=> log y

COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 7.21 215.65

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE TOTAL
ENTRY SESSION
CA SUBSCRIBER PRICE

0.00 -3.75

STN INTERNATIONAL LOGOFF AT 14:22:36 ON 20 JAN 2006